LISTING OF CLAIMS

1-14 (Cancelled).

- 15. (Previously Presented) A chimeric fusion protein comprising a bacteriorhodopsin protein amino acid sequence comprising substantially all of the amino acid sequence of bacteriorhodopsin, wherein at least a portion of the intracellular loop 3 domain of bacteriorhodopsin is replaced by at least a portion of the intracellular loop 3 domain of bootine rhodopsin, the chimeric protein having the ability to promote *in vitro* GTP-GDP exchange on transducin.
- 16. (Previously Presented) The chimeric protein of claim 15, wherein the intracellular loop 3 domain region corresponding to amino acid residues 171-179 of SEQ ID NO:2 is replaced with at least a portion of the intracellular loop 3 domain of boyine rhodopsin.
- 17. (Previously Presented) The chimeric protein of claim 16, wherein amino acid residues 171-179 of SEQ ID NO:2 are replaced with Y223-M253, Y223-R252, or Q225-R252 of bovine rhodopsin.
- (Previously Presented) A polynucleotide sequence encoding the chimeric fusion protein of claim 17.
- (Previously Presented) The polynucleotide sequence of claim 18 operably linked to a promoter.
- (Previously Presented) An archaebacterium comprising the polynucleotide sequence of claim 19.
- 21. (Previously Presented) A method of producing a bacteriorhodopsin/G protein-coupled receptor chimeric fusion protein comprising culturing the archaebacterium of claim 20 under suitable conditions and for a period of time sufficient to allow expression of the chimeric fusion protein.
- 22. (Previously Presented) A method of testing a molecule for its ability to interact with the intracellular loop 3 of a G protein-coupled receptor comprising:

- (a) contacting the chimeric fusion protein of claim 15 with the test molecule; and
- (b) detecting the presence or absence of interaction between the protein and the test molecule of step (a).
- 23. (Previously Presented) The method of claim 22 wherein the detecting step (b) comprises performing an *in vitro* GTP-GDP exchange assay.